

## DESCRIPTION

## CURABLE COMPOSITION

## Technical Field

The present invention relates to curable compositions  
5 used for adhesives, daubs, coating materials, ink, paint,  
resins for molding and the like. More particularly, the  
present invention relates to one-component epoxy curable  
compositions that can be preserved at normal temperatures.

## Background Art

10 Conventionally, many kinds of resins including epoxy  
resins, urethane resins, polysulfide resins and the like have  
been used for adhesives, architectural materials, sports goods  
and the like. However, most of such products are of a two-  
component type, wherein a resin, a curing agent and a curing  
15 accelerator are mixed immediately before use. While they are  
superior in preservation stability, they are associated with  
many defects in that an erroneous mixing ratio during use  
results in failure to cure, handling property is poor due to  
preparation when in use, which in turn increases the  
20 production cost, and the like. Thus, there is an increasing  
demand for a one-component product to omit the step for mixing  
a resin, a curing agent and a curing accelerator during use.  
To give a one-component product, for example, epoxy resins are  
used with dicyandiamide, hydrazide compounds, boron  
25 trifluoride compounds, alkylurea compounds and the like, which  
are referred to as latent curing agents, and often used  
together with a curing accelerator such as imidazole  
compounds, phosphine compounds and the like. However, such  
one-component curable composition is associated with a problem  
30 of viscosity increase as to the long-term preservation  
stability, as well as a practical problem of necessary high  
temperature heating during curing when improving the  
preservation property, and the like.

On the other hand, provision of a one-component product  
35 by the application of a microencapsulation technique has also

been studied. For example, attempts have been made to prepare a microencapsulated product of an amine curing agent by applying an interfacial polymerization method, and apply the product as a curing agent for one-component urethane resins, epoxy resins and the like (see JP-B-54-31468).

Besides these, as methods for microencapsulation of a curing agent, there have been proposed a method comprising use of a microencapsulated curing accelerator obtained by coagulating colloidal fine particles with an electrolyte to give a capsule film and covering the curing accelerator with the film (see JP-A-6-25470), a method comprising use of a latent curing agent containing at least a curing agent and one or more kinds of cyclodextrin compounds (see JP-A-9-31162) and the like.

These encapsulated curing agents are associated with problems of poor capsule stability near normal temperatures, which in turn causes viscosity increase when prepared as a one-component resin, poor reactivity upon capsule breakage, thus necessitating a long time or high temperature for curing, and the like, and practicalization thereof has been delayed.

To solve such problems, a method utilizing the solid/liquid phase change of epoxy compounds has been proposed (see JP-A-8-269167). For example, a semiconductor sealing epoxy resin composition containing a crystalline epoxy resin having a melting point of 50-150°C, a phenol resin curing agent, a curing accelerator and an inorganic filler as essential components, wherein the inorganic filler is contained in a proportion of 75-93 wt% of the whole composition, has been proposed.

It is a design where the components are mixed at a temperature lower than the crystal melting point of the epoxy resin, and in the event of curing, the composition is heated to a temperature not lower than the crystal melting point to allow the melted epoxy resin to react with the curing agent. According to this method, however, a severe reaction begins

upon melting of the epoxy resin and, particularly, a topical heat is generated due to a sharp reaction near the crystal particle interface, which in turn causes mottles during the curing reaction, and a uniformly-cured product cannot be  
5 obtained.

### Disclosure of the Invention

It is therefore an object of the present invention to provide a one-component epoxy curable composition superior in the preservation stability and having superior properties as a  
10 cured product.

The present inventors have conducted intensive studies in an attempt to solve the aforementioned problems and completed the present invention. Accordingly, the present invention provides the following.

15 (1) A curable composition that forms a continuous phase and a dispersoid at normal temperatures, wherein the continuous phase is a liquid at normal temperatures and comprises (a) a compound having two or more epoxy groups in a molecule, and

20 the dispersoid comprises (b) a compound present as solid particles in a continuous phase at normal temperatures and having two or more amino groups in a molecule.

(2) The curable composition of the above-mentioned (1), wherein the aforementioned compound having two or more amino  
25 groups in a molecule is an aromatic amine compound having a benzoxazole structure.

(3) The curable composition of the above-mentioned (1) or (2), wherein the aforementioned compound having two or more epoxy groups in a molecule is a liquid at normal temperatures.

30 (4) The curable composition of any of the above-mentioned (1)-(3), wherein the aforementioned continuous phase contains an organic solvent having a boiling point of not higher than 200°C.

(5) The curable composition of any of the above-mentioned (1)-  
35 (4), wherein the aforementioned solid particles have a volume

average particle size of 0.05 - 50  $\mu\text{m}$ .

The present invention provides a curable composition comprising an epoxy compound in a continuous phase and a multifunctional amine compound, which is a curing agent, as a solid fine particle dispersoid. The curing agent is separately present as a dispersoid from the epoxy compound in a continuous phase, and therefore, they are incompatible at least at normal temperatures, thus prohibiting a curing reaction. As a result, even though it is a one-component composition, the superior preservation stability can be maintained. However, when heated, the curing agent begins to dissolve in the continuous phase to cause reactions, which results in the production of a cured product.

When the change of solid/liquid phases of a compound is utilized, which has been conventionally proposed as to the use of a crystalline epoxy resin, an epoxy compound activated due to high temperature reaching the melting point is suddenly brought into contact with a curing agent. Thus, a severe reaction occurs and mottles are generated in the cured product. As shown in the present invention, when the both are brought into contact with each other through dissolution, the reaction occurs comparatively gently, the epoxy compound and the curing agent are mixed comparatively uniformly, the structure of the cured product becomes uniform and, as a result, superior properties of the cured product can be achieved.

#### **Detailed Description of the Invention**

The curable composition of the present invention forms, at normal temperatures, a continuous phase, which is a liquid at normal temperatures, comprising (a) a compound having two or more epoxy groups in a molecule (hereinafter sometimes to be abbreviated as epoxy compound (a)) and, as a curing agent, a dispersoid comprising (b) a compound present as solid particles in a continuous phase at normal temperatures and having two or more amino groups in a molecule (hereinafter

sometimes to be abbreviated as amine compound (b)). Due to such constitution, a curing reaction does not occur, since epoxy compound (a) and amine compound (b) are incompatible at least at normal temperature, and superior preservation  
5 stability can be maintained. In the present invention, the normal temperature means the surrounding temperature in the environment, and generally means a temperature within the range of about -20°C to 50°C.

As epoxy compound (a) in the present invention, a  
10 compound having two or more epoxy groups in one molecule can be used. As such compound, glycidyl ether epoxy compounds, such as glycidyl ether of bisphenol A, glycidyl ether of bisphenol F, glycidyl ether of bisphenol S, glycidyl ether of resorcin, glycidyl ether of glycerol, glycidyl ether of  
15 polyalkylene oxide, glycidyl ether of brominated bisphenol A and oligomers thereof, and the like can be mentioned. Furthermore, an epoxy compound which is a condensate of phenols, naphthols and the like with formalins, aliphatic or aromatic aldehydes, or ketones, as represented by glycidyl  
20 ether of phenol novolac, glycidyl ether of cresol novolac and the like, an alicyclic epoxy compound, such as alicyclic diepoxyacetal, alicyclic diepoxy adipate, alicyclic diepoxy carboxylate and the like can be mentioned.

In addition, glycidyl ester epoxy compounds (i.e.  
25 phthalic acid diglycidyl ester, tetrahydrophthalic acid diglycidyl ester, hexahydrophthalic acid diglycidyl ester and the like), glycidyl amine epoxy compounds (i.e. N,N-diglycidylaniline, tetraglycidylaminodiphenylmethane), heterocyclic epoxy compound, hydantoin epoxy compound,  
30 triglycidyl isocyanurate and the like can be mentioned. These epoxy compounds (a) can be used alone, or in a combination of two or more kinds thereof.

In the present invention, use of epoxy compound (a) having two or more epoxy groups in a molecule is essential. In  
35 addition to such compound, a monofunctional epoxy compound may



be used in combination as necessary.

The continuous phase is a liquid at normal temperatures. The epoxy compound (a) preferably used to afford a liquid continuous phase is an epoxy compound having a comparatively  
5 low molecular weight, which is a liquid at normal temperatures, such as glycidyl ether of bisphenol A, glycidyl ether of phenol novolac and the like. An epoxy compound (a) having a molecular weight of not more than 1000, preferably not more than 800, more preferably not more than 600, and  
10 still more preferably not more than 400, can be preferably used. The epoxy compound (a) does not need to be a liquid in itself and, for example, an epoxy compound having a comparatively high molecular weight, which is a solid in itself, may be dissolved in a low molecular weight liquid  
15 epoxy compound.

In the present invention, a liquid continuous phase may be afforded by adding a conventional low-boiling solvent to a continuous phase. The combined use of a solvent is effective when epoxy compound (a) is a solid or has an extremely high  
20 viscosity. Moreover, the use of a solvent is preferable when a coated film having a small thickness is desired in the application to daub or adhesive. As the organic solvent to be used, a solvent having a boiling point of not more than 200°C is preferable, a solvent having a boiling point of not more  
25 than 180°C is more preferable and a solvent having a boiling point of not more than 160°C is still more preferable. The ultimate object of the present invention is provision of a cured product. Therefore, the use of a high-boiling solvent is not preferable, since it degrades the properties of the cured  
30 product. Moreover, when the boiling point of the solvent is too low, practicality is impaired. Thus, a solvent having a boiling point of not less than 50°C is preferable, a solvent having a boiling point of not less than 70°C is more preferable, and a solvent having a boiling point of not less  
35 than 90°C is still more preferable.

More concrete solvents are selected from, for example, methanol, ethanol, propanol, butanol, toluene, xylene, Solvesso, Isopar, ethyl acetate, butyl acetate, methyl ethyl ketone, cyclohexanone, cyclohexanol, isophorone, methyl  
5 cellosolve, ethyl cellosolve, butyl cellosolve, butyl cellosolve acetate, ethyl Carbitol, ethyl Carbitol acetate, ethyleneglycolmonoacetate, dimethylformamide,  $\gamma$ -butyrolactone, n-methyl pyrrolidone and the like, in consideration of solubility, evaporation rate and the like. While the amount of  
10 the solvent to be added varies depending on the epoxy compound (a) to be used, it is generally 10-1000 parts by weight, preferably 30-300 parts by weight, per 100 parts by weight of epoxy compound (a). When the amount of the solvent is too high, the viscosity of a curable composition decreases to  
15 unpreferably make uniform coating difficult.

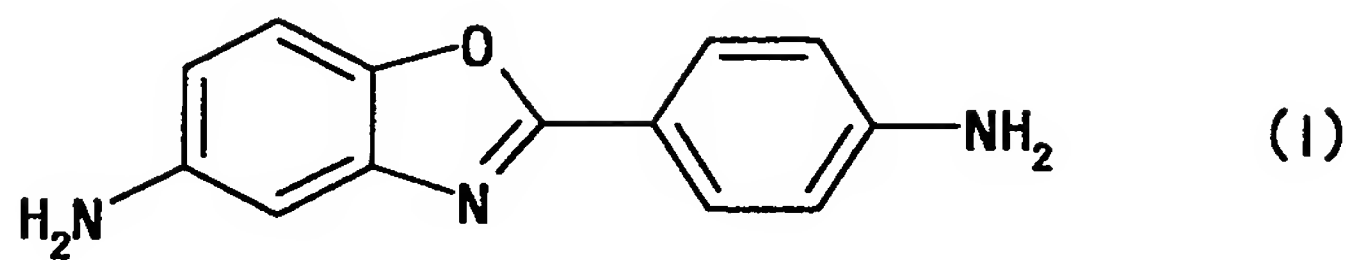
In the present invention, use of amine compound (b) as an curing agent is essential, which has two or more amino groups in a molecule and is dispersed as solid particles in a continuous phase containing epoxy compound (a) at normal  
20 temperatures.

The melting point of amine compound (b) is preferably 60-400°C, more preferably 100-350°C. When the melting point is too low, the compound is likely to elute out to impair preservation stability, and when it is too high, curing is  
25 likely to require high temperature and long time, thus degrading the operability.

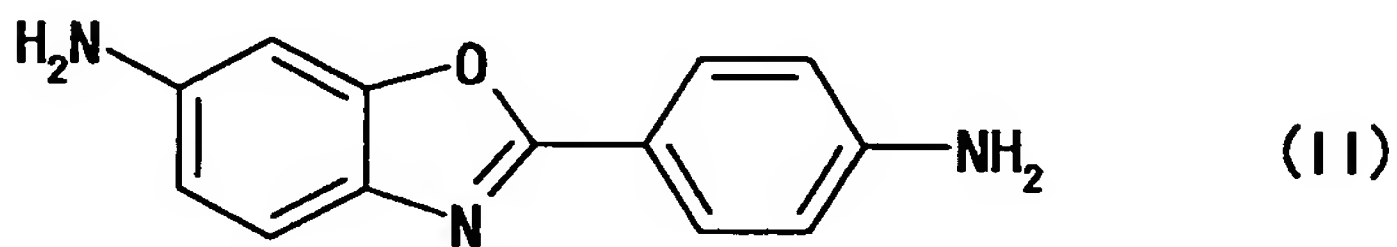
As amine compound (b) satisfying such conditions, for example, a multifunctional amine compound having the aromatic heterocyclic skeleton can be preferably mentioned. Moreover, a  
30 multifunctional amine compound having the aromatic heteronuclear skeleton can be preferably mentioned.

In the present invention, an aromatic amine compound having a benzoxazole structure can be preferably used as amine compound (b). As the aromatic amine compound having a  
35 benzoxazole structure, compounds represented by the following

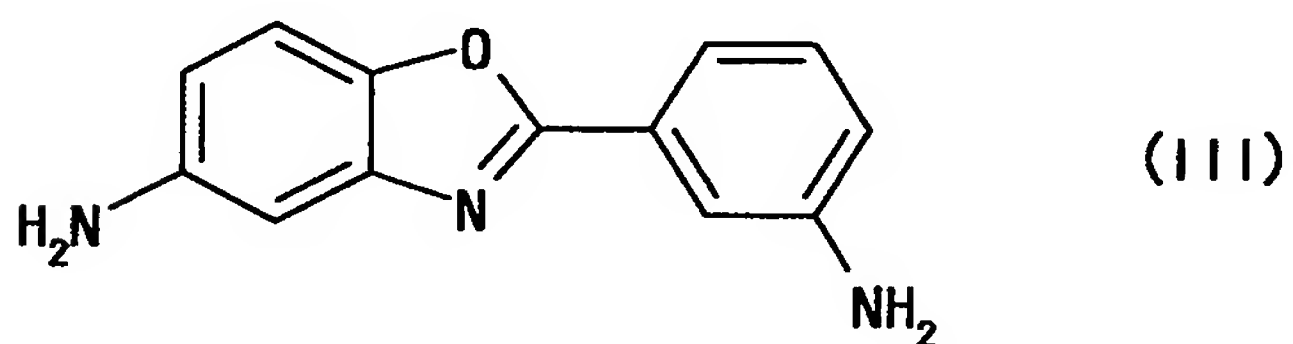
chemical formulas (I)-(XIII) can be mentioned.



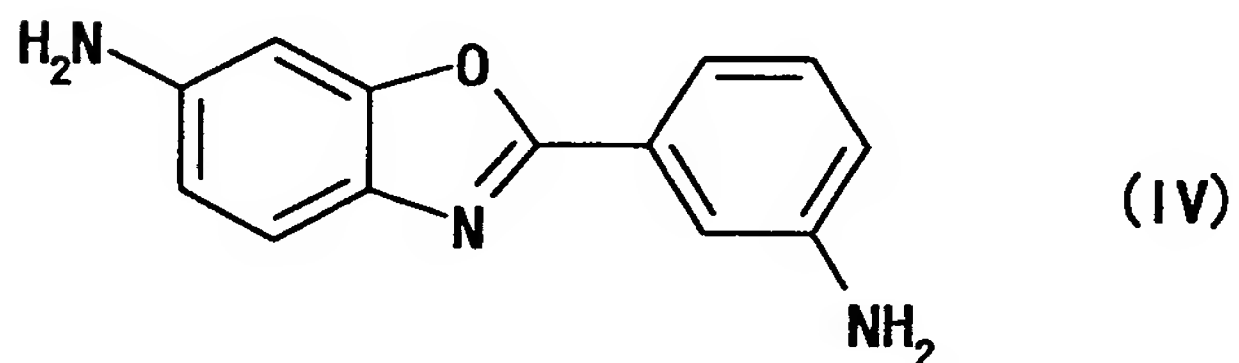
5-amino-2-(p-aminophenyl)benzoxazole



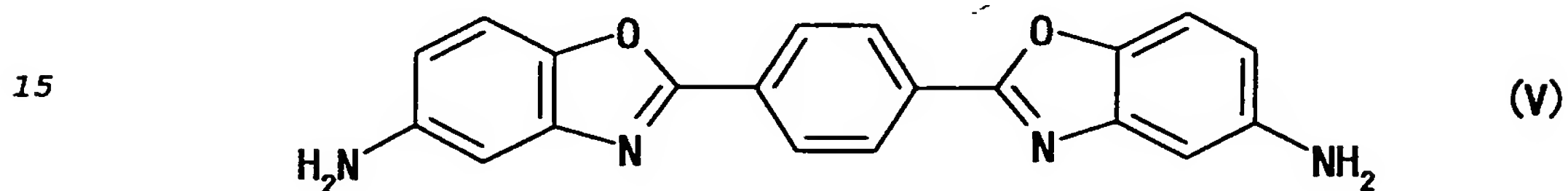
6-amino-2-(p-aminophenyl)benzoxazole



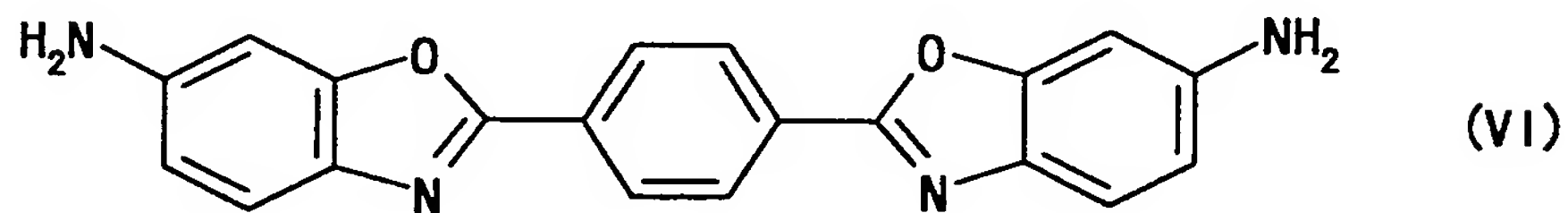
5-amino-2-(m-aminophenyl)benzoxazole



6-amino-2-(m-aminophenyl)benzoxazole

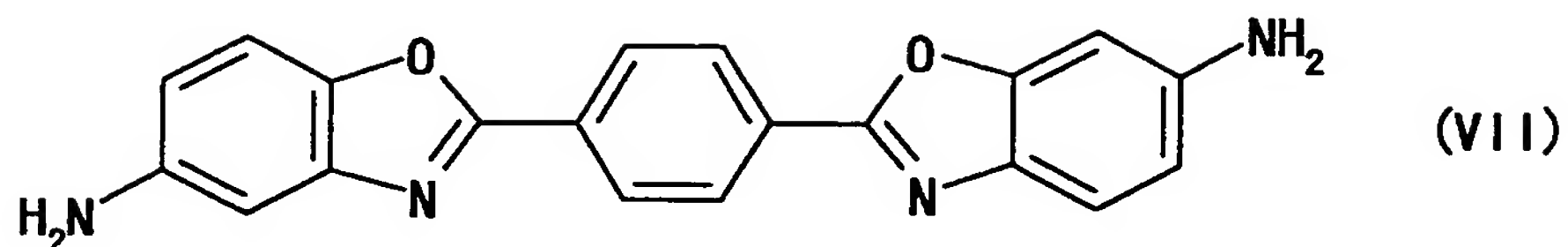


2,2'-p-phenylenebis(5-aminobenzoxazole)

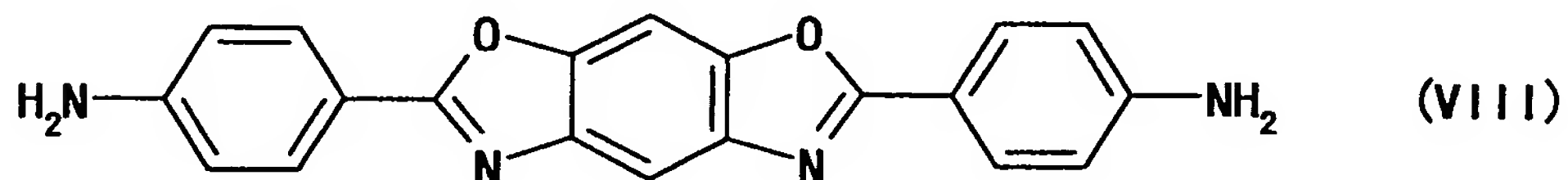


2,2'-p-phenylenebis(6-aminobenzoxazole)

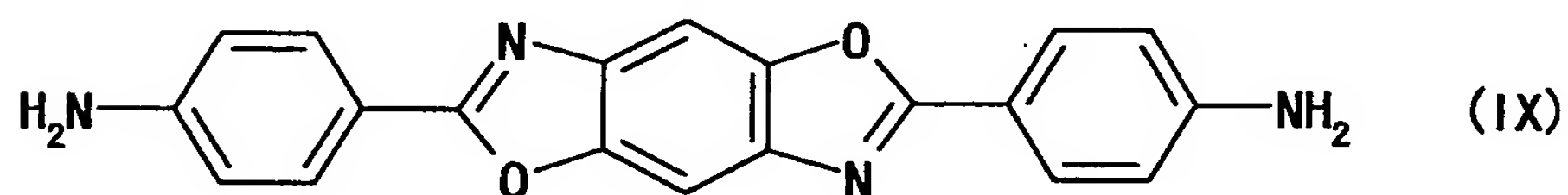




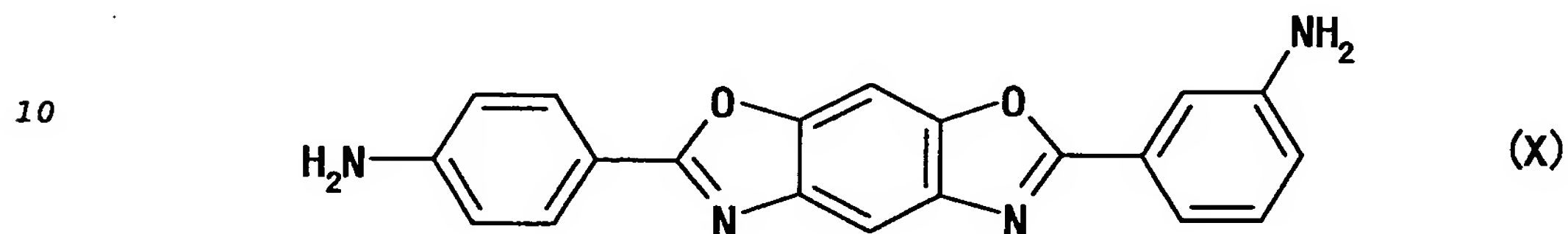
1-(5-aminobenzoxazolo)-4-(6-aminobenzoxazolo)benzene



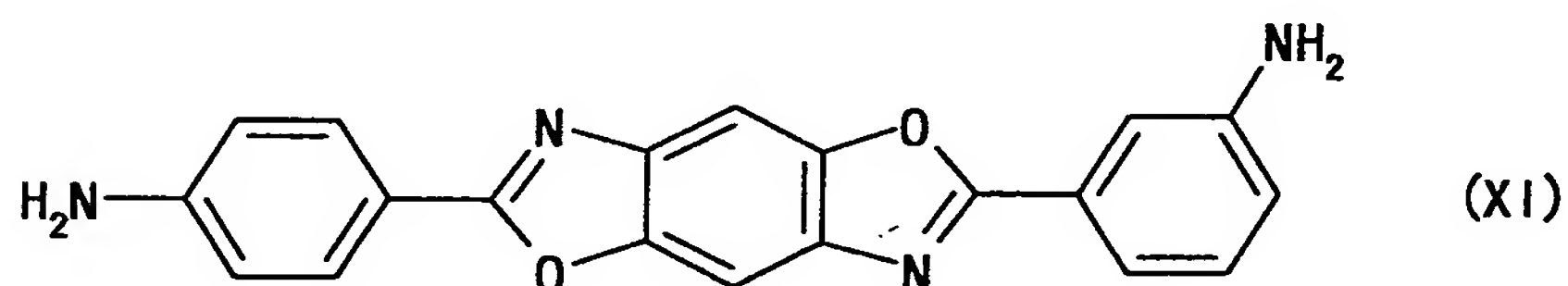
5 2,6-(4,4'-diaminodiphenyl)benzo[1,2-d:5,4-d']bisoxazole



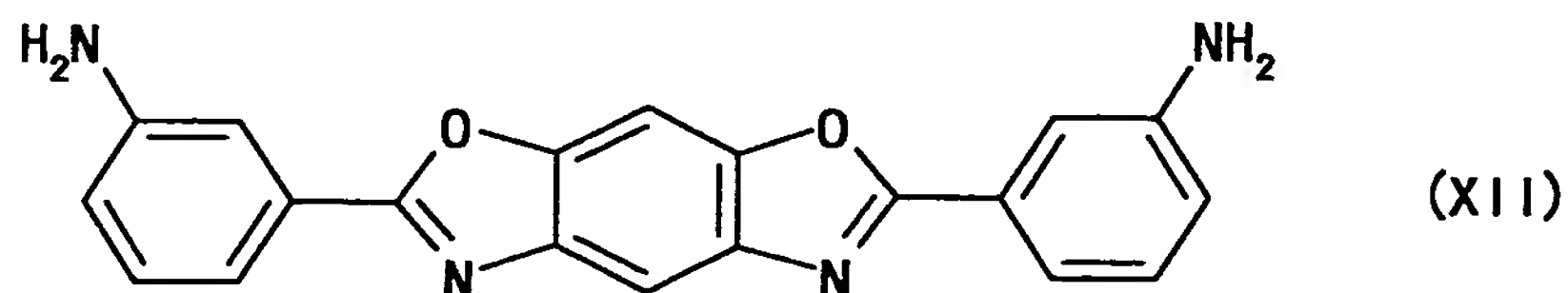
2,6-(4,4'-diaminodiphenyl)benzo[1,2-d:4,5-d']bisoxazole



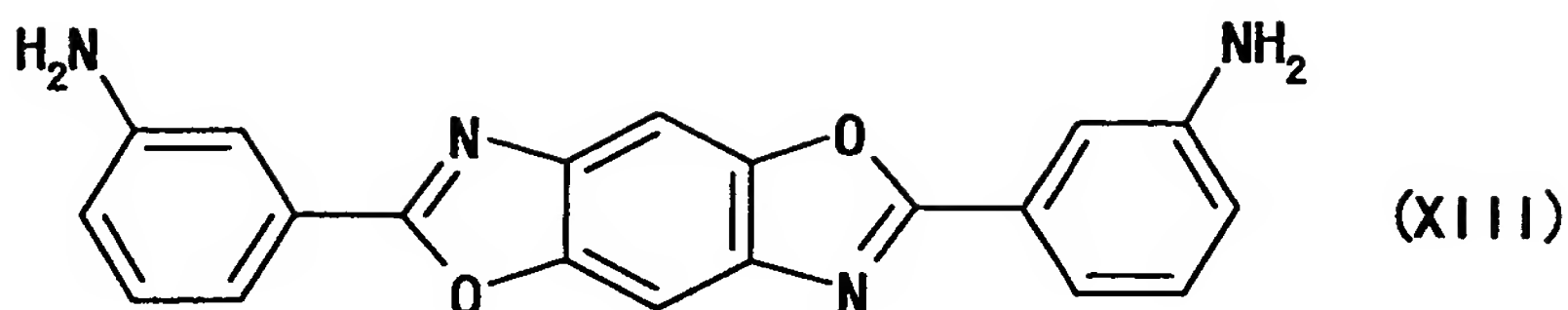
10 2,6-(3,4'-diaminodiphenyl)benzo[1,2-d:5,4-d']bisoxazole



15 2,6-(3,4'-diaminodiphenyl)benzo[1,2-d:4,5-d']bisoxazole



2,6-(3,3'-diaminodiphenyl)benzo[1,2-d:5,4-d']bisoxazole



2,6-(3,3'-diaminodiphenyl)benzo[1,2-d:4,5-d']bisoxazole

The amine compounds (b) may be used alone or in a  
 5 mixture of two or more kinds thereof.

In the present invention, an amine compound incapable of existing in the form of solid particles in a continuous phase due to dissolution in a continuous phase at normal temperatures, for example, one or more kinds of diamines free  
 10 of a benzoxazole structure, which are exemplified in the following, may be used in combination as long as the preservation stability is not impaired, which is generally not more than 30 mol%, preferably not more than 15 mol%, of the total diamine. As such diamines, for example, 4,4'-bis(3-  
 15 aminophenoxy)biphenyl, bis[4-(3-aminophenoxy)phenyl]ketone, bis[4-(3-aminophenoxy)phenyl]sulfide, bis[4-(3-aminophenoxy)phenyl]sulfone, 2,2-bis[4-(3-aminophenoxy)phenyl]propane, 2,2-bis[4-(3-aminophenoxy)phenyl]-1,1,1,3,3,3-hexafluoropropane, m-  
 20 phenylenediamine, o-phenylenediamine, p-phenylenediamine, m-aminobenzylamine, p-aminobenzylamine, 3,3'-diaminodiphenylether, 3,4'-diaminodiphenylether, 4,4'-diaminodiphenylether, 3,3'-diaminodiphenylsulfide, 3,3'-diaminodiphenylsulfoxide, 3,4'-diaminodiphenylsulfoxide, 4,4'-  
 25 diaminodiphenylsulfoxide, 3,3'-diaminodiphenylsulfone, 3,4'-diaminodiphenylsulfone, 4,4'-diaminodiphenylsulfone, 3,3'-diaminobenzophenone, 3,4'-diaminobenzophenone, 4,4'-diaminobenzophenone, 3,3'-diaminodiphenylmethane, 3,4'-diaminodiphenylmethane, 4,4'-diaminodiphenylmethane, bis[4-(4-  
 30 aminophenoxy)phenyl]methane, 1,1-bis[4-(4-aminophenoxy)phenyl]ethane, 1,2-bis[4-(4-aminophenoxy)phenyl]ethane, 1,1-bis[4-(4-

aminophenoxy)phenyl]propane, 1,2-bis[4-(4-  
 aminophenoxy)phenyl]propane, 1,3-bis[4-(4-  
 aminophenoxy)phenyl]propane, 2,2-bis[4-(4-  
 aminophenoxy)phenyl]propane, 1,1-bis[4-(4-  
 5 aminophenoxy)phenyl]butane, 1,3-bis[4-(4-  
 aminophenoxy)phenyl]butane, 1,4-bis[4-(4-  
 aminophenoxy)phenyl]butane, 2,2-bis[4-(4-  
 aminophenoxy)phenyl]butane, 2,3-bis[4-(4-  
 aminophenoxy)phenyl]butane, 2-[4-(4-aminophenoxy)phenyl]-2-[4-  
 10 (4-aminophenoxy)-3-methylphenyl]propane, 2,2-bis[4-(4-  
 aminophenoxy)-3-methylphenyl]propane, 2-[4-(4-  
 aminophenoxy)phenyl]-2-[4-(4-aminophenoxy)-3,5-  
 dimethylphenyl]propane, 2,2-bis[4-(4-aminophenoxy)-3,5-  
 dimethylphenyl]propane, 2,2-bis[4-(4-aminophenoxy)phenyl]-  
 15 1,1,1,3,3,3-hexafluoropropane, 1,4-bis(3-aminophenoxy)benzene,  
 1,3-bis(3-aminophenoxy)benzene, 1,4-bis(4-  
 aminophenoxy)benzene, 4,4'-bis(4-aminophenoxy)biphenyl, bis[4-  
 (4-aminophenoxy)phenyl]ketone, bis[4-(4-  
 aminophenoxy)phenyl]sulfide, bis[4-(4-  
 20 aminophenoxy)phenyl]sulfoxide, bis[4-(4-  
 aminophenoxy)phenyl]sulfone, bis[4-(3-  
 aminophenoxy)phenyl]ether, bis[4-(4-aminophenoxy)phenyl]ether,  
 1,3-bis[4-(4-aminophenoxy)benzoyl]benzene, 1,3-bis[4-(3-  
 aminophenoxy)benzoyl]benzene, 1,4-bis[4-(3-  
 25 aminophenoxy)benzoyl]benzene, 4,4'-bis(3-  
 aminophenoxy)benzoyl]benzene, 1,1-bis[4-(3-  
 aminophenoxy)phenyl]propane, 1,3-bis[4-(3-  
 aminophenoxy)phenyl]propane, 3,4'-diaminodiphenylsulfide, 2,2-  
 bis[3-(3-aminophenoxy)phenyl]-1,1,1,3,3,3-hexafluoropropane,  
 30 bis[4-(3-aminophenoxy)phenyl]methane, 1,1-bis[4-(3-  
 aminophenoxy)phenyl]ethane, 1,2-bis[4-(3-  
 aminophenoxy)phenyl]ethane, bis[4-(3-  
 aminophenoxy)phenyl]sulfoxide, 4,4'-bis[3-(4-  
 aminophenoxy)benzoyl]diphenyl ether, 4,4'-bis[3-(3-  
 35 aminophenoxy)benzoyl]diphenyl ether, 4,4'-bis[4-(4-amino- $\alpha,\alpha$ -

dimethylbenzyl)phenoxy]benzophenone, 4,4'-bis[4-(4-amino- $\alpha,\alpha$ -  
 dimethylbenzyl)phenoxy]diphenylsulfone, bis[4-{4-(4-  
 aminophenoxy)phenoxy}phenyl]sulfone, 1,4-bis[4-(4-  
 aminophenoxy)phenoxy- $\alpha,\alpha$ -dimethylbenzyl]benzene, 1,3-bis[4-(4-  
 5 aminophenoxy)phenoxy- $\alpha,\alpha$ -dimethylbenzyl]benzene, 1,3-bis[4-(4-  
 amino-6-trifluoromethylphenoxy)- $\alpha,\alpha$ -dimethylbenzyl]benzene,  
 1,3-bis[4-(4-amino-6-fluorophenoxy)- $\alpha,\alpha$ -  
 dimethylbenzyl]benzene, 1,3-bis[4-(4-amino-6-methylphenoxy)-  
 $\alpha,\alpha$ -dimethylbenzyl]benzene, 1,3-bis[4-(4-amino-6-  
 10 cyanophenoxy)- $\alpha,\alpha$ -dimethylbenzyl]benzene, 3,3'-diamino-4,4'-  
 diphenoxybenzophenone, 4,4'-diamino-5,5'-  
 diphenoxybenzophenone, 3,4'-diamino-4,5'-  
 diphenoxybenzophenone, 3,3'-diamino-4-phenoxybenzophenone,  
 4,4'-diamino-5-phenoxybenzophenone, 3,4'-diamino-4-  
 15 phenoxybenzophenone, 3,4'-diamino-5'-phenoxybenzophenone,  
 3,3'-diamino-4,4'-dibiphenoxybenzophenone, 4,4'-diamino-5,5'-  
 dibiphenoxybenzophenone, 3,4'-diamino-4,5'-  
 dibiphenoxybenzophenone, 3,3'-diamino-4-biphenoxybenzophenone,  
 4,4'-diamino-5-biphenoxybenzophenone, 3,4'-diamino-4-  
 20 biphenoxybenzophenone, 3,4'-diamino-5'-biphenoxybenzophenone,  
 1,3-bis(3-amino-4-phenoxybenzoyl)benzene, 1,4-bis(3-amino-4-  
 phenoxybenzoyl)benzene, 1,3-bis(4-amino-5-  
 phenoxybenzoyl)benzene, 1,4-bis(4-amino-5-  
 phenoxybenzoyl)benzene, 1,3-bis(3-amino-4-  
 25 biphenoxybenzoyl)benzene, 1,4-bis(3-amino-4-  
 biphenoxybenzoyl)benzene, 1,3-bis(4-amino-5-  
 biphenoxybenzoyl)benzene, 1,4-bis(4-amino-5-  
 biphenoxybenzoyl)benzene, 2,6-bis[4-(4-amino- $\alpha,\alpha$ -  
 dimethylbenzyl)phenoxy]benzonitrile and the above-mentioned  
 30 aromatic diamines wherein the hydrogen atoms on the aromatic  
 ring are partially or entirely substituted by a halogen atom,  
 an alkyl group or alkoxy group having 1 to 3 carbon atoms, a  
 cyano group, or a halogenated alkyl group or alkoxy group  
 having 1 to 3 carbon atoms, wherein the hydrogen atoms of the  
 35 alkyl group or alkoxy group are partially or entirely

substituted by a halogen atom, and the like can be mentioned.

In the present invention, amine compound (b), preferably an aromatic diamine compound having a benzoxazole structure is dispersed in the composition in a solid fine particle state.

5 In this case, its volume average particle size is preferably 0.05-50  $\mu\text{m}$ , more preferably 0.1-15  $\mu\text{m}$ , and still more preferably 0.3-7  $\mu\text{m}$ . When the volume average particle size exceeds a given range, the composition is likely to become non-uniform due to the precipitation or floatation of the

10 dispersed particles to result in a mottled cured film. When it is smaller than the given range, the viscosity of the whole system is likely to increase markedly, thus making the handling difficult.

In the present invention, the volume average particle

15 size means a volume average particle size calculated from the particle size distribution measured by the laser scattering method.

To be specific, the curable composition of the present invention is diluted 100-fold (v/v) with toluene, the particle

20 size distribution is measured using a laser scattering particle size distribution analyzer (e.g., LB-500 manufactured by Horiba, Ltd., and the like), and the volume average particle size is calculated with the refractive index (1.496) of toluene used as a refractive index ( $n_m$ ) of the medium and

25 1.55 as the refractive index ( $n_p$ ) of dispersoid.

The particle size distribution of amine compound (b) is preferably sharp. To be specific, the standard deviation/average value (CV value) measured according to the laser scattering method is preferably not more than 1.0, more

30 preferably not more than 0.7, and still more preferably not more than 0.5.

In the present invention, amine compound (b) can be used by dividing ultrafinely in a dry mill such as a chopper mill, a jet mill, an angmill, an atomizer and the like.

35 Alternatively, it can be dispersed ultrafinely during



dispersing using a wet-dispersing device such as an attritor, a homogenizer, a nanomizer, and the like. It is also possible to concurrently use a known dispersant to the extent the properties of the cured product is not impaired.

5       The mixing ratio of the epoxy compound (a) and amine compound (b) in the present invention, A/B, wherein the total molar amount of epoxy group in the epoxy compound (a) is A (mol) and the total molar amount of amino group in the amine compound (b) is B (mol), is preferably 0.70-1.50, more  
10 preferably 0.80-1.35, still more preferably 0.90-1.2, and further preferably 0.95-1.08. When the mixing ratio exceeds a given range, the properties of the cured product are markedly degraded. When the mixing ratio is less than a given range, the properties of the cured product are degraded, the water  
15 absorbability of the cured product increases, and the permanence and electric insulation are degraded.

      In the present invention, an appropriate amount of a conventionally used curing agent may be concurrently used as the curing agent. As such curing agent, amine curing agents,  
20 acid anhydride curing agents, phenol curing agents and the like can be mentioned. To be specific, aliphatic diamines, aliphatic polyamines, aliphatic polyamines including aromatic ring, alicyclic and cyclic polyamines, aromatic primary amines etc., aliphatic acid anhydrides, alicyclic acid anhydrides,  
25 aromatic acid anhydrides, halogen acid anhydrides, trisphenol, phenol novolac, cresol novolac and the like can be mentioned. Since the concurrent use of these curing agents may impair the preservation stability, its amount of addition is not more than 10 wt%, preferably not more than 5 wt%, more preferably  
30 not more than 2 wt%, relative to the total amount of the curable composition.

      In the present invention, a curing accelerator may be used as necessary. As the curing accelerator, 1,8-diaza-bicyclo-(5,4,0)-undecene-7 (DBU), phosphorus curing  
35 accelerators (e.g., tributylphosphine, trioctylphosphine,

tricyclohexylphosphine, triphenylphosphine (TPP),  
tribenzylphosphine, tritolylphosphine, p-styrylphosphine,  
tris(2,6-dimethoxyphenyl)phosphine, tri-4-  
methylphenylphosphine, tri-2-cyanoethylphosphine and the like,  
5 bis(diphenylphosphino)methane, 1,2-  
bis(diphenylphosphino)ethane, 1,4-(diphenylphosphino)butane,  
triphenylphosphine-triphenylborane, tetraphenylphosphonium  
tetraphenylborate and the like), imidazole curing accelerators  
(e.g., 2-methylimidazole, 2-ethyl-4-methylimidazole, 2-phenyl-  
10 4-methylimidazole, 1-benzyl-2-methylimidazole, 2-  
ethylimidazole, 2-isopropylimidazole, 2-undecylimidazole, 1-  
cyanoethyl-2-methylimidazole, 1-cyanoethyl-2-ethyl-4-  
methylimidazole, 1-cyanoethyl-2-undecylimidazole, 1-  
cyanoethyl-2-isopropylimidazole, 1-cyanoethyl-2-  
15 phenylimidazole), and the like can be mentioned. The curing  
accelerator is not limited to these and various curing  
accelerators can be used. These curing accelerators can be  
used alone or in a mixture thereof as necessary. The amount of  
the curing accelerator to be added can be appropriately  
20 selected from the range of 0.1-10 wt% relative to the total  
amount of the curable composition, so that an appropriate  
curing performance can be obtained in consideration of the  
epoxy compound (a) and amine compound (b) to be used.

In the present invention, moreover, dye, inorganic  
25 filler, flexibilizer, organic filler, solvent, diluent,  
pigment, flame-retardant, mold lubricant, silane coupling  
agent, titanium coupling agent and the like can be added as  
necessary, as long as the action effect of the present  
invention is not impaired.

30 The curable composition of the present invention can be  
obtained by adding a dispersoid to a continuous phase  
containing epoxy compound (a) and finely dispersing the  
dispersoid. For fine dispersion of the materials, known  
dispersion means such as a sand mill, an attritor, a shaker, a  
35 dissolver, a ball mill, a roll mill and the like can be used.

In the curable composition of the present invention, heating causes dissolution and elution of amine compound (b) in a continuous phase, and progress of the curing reaction. While the heating temperature varies depending on the epoxy compound (a) and amine compound (b) to be used, it is generally within the range of 80-250°C, preferably 100-200°C, and the curing can be carried out in 0.1-6 hr. When the heating temperature is too high, the cured product is likely to become non-uniform or deteriorated. When it is too low, the curing degree of the cured product is likely to be insufficient.

### Examples

The present invention is explained in detail in the following by referring to Examples. In the Examples and Comparative Examples, the physical values were evaluated by the following method.

#### (1) Volume average particle size

A curable composition was diluted 100-fold with toluene, placed in a quartz cell, and the volume average particle size was measured by a laser scattering particle size distribution analyzer LB-500 manufactured by Horiba, Ltd. It was calculated using the values of toluene as the property values and the refractive index of the medium and 1.55 as the refractive index of the dispersoid.

#### (2) Preservation stability (viscosity increase)

A curable composition was stood at 50°C for 1000 hr, and the gel fraction was measured and the preservation stability was evaluated. The evaluation criteria were as follows.

- O: gel fraction  $\leq 5\%$
- $\Delta$ :  $5\% < \text{gel fraction} \leq 10\%$
- x:  $10\% < \text{gel fraction}$

The gel fraction was determined by Soxhlet extracting the soluble matters from the cured product for 24 hr with N-methylpyrrolidone as a solvent, and from the following

formula.

Gel fraction (%) = (weight of sample after extraction and drying) / (weight of sample before extraction) × 100

### (3) Curability

5 A curable composition was stood at 150°C for 1 hr, and the gel fraction was measured and the curability was determined. The evaluation criteria were as follows.

O: 90% ≤ gel fraction

Δ: 80% ≤ gel fraction < 90%

10 x: gel fraction < 80%

The gel fraction was determined by Soxhlet extracting the soluble matters from the cured product for 24 hr with N-methylpyrrolidone as a solvent, and from the following formula.

15 Gel fraction (%) = (weight of sample after extraction and drying) / (weight of sample before extraction) × 100

### (4) Pencil hardness of cured product

Pencil hardness was measured according to JIS-K5400.

### (5) Adhesive property of cured product

20 A curable composition was laminated on a polyimide film KAPTON 100H (polyimide film, manufactured by DU PONT-TORAY CO. LTD.) and the 90-degree peel strength was measured after heat hardening at 150°C for 2 hr.

### (6) Water resistance of cured product

25 A curable composition was immersed in boiling water at 100°C, left for standing for 2 hr, taken out and the appearance of the cured product was observed. Water resistance was evaluated by marking those without change in the appearance after the treatment with O, and those with crack or whitening  
30 with x.

### (Example 1)

Epikote 828 (19.0 parts by weight, epoxy equivalent amount 190 g/eq., bisphenol A epoxy compound manufactured by Japan Epoxy Resins Co. Ltd.) and 5-amino-2-(p-  
35 aminophenyl)benzoxazole (11.3 parts by weight, compound of the

formula (I), amine equivalent amount 113 g/eq.) were charged in a 100 ml flask, and the mixture was stirred at room temperature for 5 min and dispersed with a three roll mill to give a curable composition, which was subjected to each  
5 evaluation test. This curable composition was heated at 150°C for 2 hr to give a cured product.

(Example 2)

In the same manner as in Example 1 and using Epikote 828 (19.0 parts by weight, epoxy equivalent amount 190 g/eq.,  
10 bisphenol A epoxy compound manufactured by Japan Epoxy Resins Co. Ltd.) and 2,2'-p-phenylenebis(5-aminobenzoxazole)benzoxazole (17.1 parts by weight, compound of the formula (V), amine equivalent amount 171 parts by weight/eq.), a curable composition was obtained and subjected  
15 to the evaluation in the same manner.

(Example 3)

In the same manner as in Example 1 and using Epikote 828 (19.0 parts by weight, epoxy equivalent amount 190 g/eq.) and 2,6-(4,4'-diaminodiphenyl)benzo[1,2-d:4,5-d']bisoxazole (17.1  
20 parts by weight, compound of the formula (IX), amine equivalent amount 171 g/eq.), a curable composition was obtained and subjected to the evaluation in the same manner.

(Example 4)

In the same manner as in Example 1 and using EOCN-102S  
25 (21.0 parts by weight, epoxy equivalent amount 210 g/eq., cresol novolac epoxy compound manufactured by NIHON KAYAKU CO. LTD.) and 5-amino-2-(p-aminophenyl)benzoxazole (11.3 parts by weight, compound of the formula (I), amine equivalent amount 113 g/eq.), a curable composition was obtained and subjected  
30 to the evaluation in the same manner.

(Example 5)

Epikote 1001 (47.5 parts by weight, 0.050 mol, epoxy equivalent amount 475 g/eq, bisphenol A epoxy compound manufactured by Japan Epoxy Resins Co. Ltd.) was dissolved in  
35 methylethylketone (100 parts by weight), 5-amino-2-(p-



aminophenyl)benzoxazole (11.3 parts by weight, compound of the formula (I), amine equivalent amount 113 g/eq.) was added, and the mixture was stirred at room temperature for 5 min and dispersed for 3 hrs in a ball mill with an alumina ball  
5 (diameter 10 mm) to give a curable composition, which was subjected to each evaluation test. This curable composition was applied to a given substrate, hot-air dried at 80°C for 30 min and heated at 150°C for 2 hrs to give a cured product.

(Example 6)

10 In the same manner as in Example 5 and using Epikote 1004 (92.5 parts by weight, epoxy equivalent amount 925 g/eq., bisphenol A epoxy compound manufactured by Japan Epoxy Resins Co. Ltd.), 5-amino-2-(p-aminophenyl)benzoxazole (11.3 parts by weight, compound of the formula (I), amine equivalent amount  
15 113 g/eq.) and methylethylketone (200 parts by weight), a curable composition was obtained and subjected to the evaluation in the same manner.

(Comparative Example 1)

In the same manner as in Example 1 and using Epikote 828  
20 (19.0 parts by weight, epoxy equivalent amount 190 g/eq.) and o-phenylenediamine (5.4 parts by weight, amine equivalent amount 54 g/eq), a curable composition was obtained and subjected to the evaluation in the same manner. o-Phenylenediamine became compatible with Epikote 828 by  
25 stirring at room temperature for 5 min. For confirmation, the volume average particle size was measured. As a result, it was smaller than the measurable level of 0.05  $\mu\text{m}$ , and the absence of substantially solid particles was confirmed. The curable composition showed remarkable viscosity increase from  
30 immediately after preparation to the extent that the composition could not be taken out from the container after standing overnight.

(Comparative Example 2)

In the same manner as in Example 1 and using Epikote 828  
35 (19.0 parts by weight, epoxy equivalent amount 190 g/eq.) and

m-phenylenediamine (5.4 parts by weight, amine equivalent amount 54 g/eq), a curable composition was obtained and subjected to the evaluation in the same manner. m-Phenylenediamine became compatible with Epikote 828 by stirring at room temperature for 5 min. For confirmation, the volume average particle size was measured. As a result, it was smaller than the measurable level of 0.05  $\mu\text{m}$ , and the absence of substantially solid particles was confirmed. Like Comparative Example 1, the curable composition showed remarkable viscosity increase from immediately after preparation to the extent that the composition could not be taken out from the container after standing overnight. (Comparative Example 3)

In the same manner as in Example 1 and using Epikote 828 (19.0 parts by weight, epoxy equivalent amount 190 g/eq.) and p-phenylenediamine (5.4 parts by weight, amine equivalent amount 54 g/eq), a curable composition was obtained and subjected to the evaluation in the same manner. p-Phenylenediamine became compatible with Epikote 828 by stirring at room temperature for 5 min. For confirmation, the volume average particle size was measured. As a result, it was smaller than the measurable level of 0.05  $\mu\text{m}$ , and the absence of substantially solid particles was confirmed. Like Comparative Example 1, the curable composition showed remarkable viscosity increase from immediately after preparation to the extent that the composition could not be taken out from the container after standing overnight. (Comparative Example 4)

In the same manner as in Example 1 and using Epikote 828 (19.0 parts by weight, epoxy equivalent amount 190 g/eq.) and ethylenediamine (3.0 parts by weight, amine equivalent amount 30 g/eq), a curable composition was obtained and subjected to the evaluation in the same manner. Ethylenediamine became compatible with Epikote 828 by stirring at room temperature for 5 min. An exothermic curing reaction occurred from

immediately after mixing and the usable time was of the order of several minutes.

(Comparative Example 5)

In the same manner as in Example 1 and using Epikote 828  
5 (19.0 parts by weight, epoxy equivalent amount 190 g/eq.) and  
diaminodiphenyl ether (16.6 parts by weight, amine equivalent  
amount 166 g/eq), a curable composition was obtained and  
subjected to the evaluation in the same manner.

Diaminodiphenyl ether became compatible with Epikote 828 by  
10 stirring at room temperature for 5 min. For confirmation, the  
volume average particle size was measured. As a result, it was  
smaller than the measurable level of 0.05  $\mu\text{m}$ , and the absence  
of substantially solid particles was confirmed. An exothermic  
curing reaction occurred from immediately after mixing and the  
15 usable time was of the order of several minutes.

(Comparative Example 6): microencapsulated curing accelerator

DBU: 1,8-diaza-bicyclo-(5,4,0)-undecene-7 (80 parts by  
weight) was dissolved in silica sol (2000 parts by weight,  
SNOWTEX O, manufactured by NISSAN CHEMICAL INDUSTRIES, LTD.,  
20 particle size 10-20 nm, concentration 20%), and the solution  
was dispersed in a solution of sorbitan polyoxyethylene  
monooleate (8 parts by weight) and sorbitan trioleate (80  
parts by weight) in chloroform (6000 parts by weight) with a  
homogenizer (8000 rpm, 30 sec) to give a W/O dispersion. This  
25 was placed in a round flask with a stirrer, and 20 wt% aqueous  
calcium chloride solution (1000 parts by weight) was added  
dropwise over about 5 min at room temperature with stirring.  
The stirring was continued for 15-30 min at room temperature.  
To a slurry of the obtained capsule particles was added  
30 methanol (1000-2000 parts by weight), capsule particles were  
filtered off by suction filtration and dried to give a  
microencapsulated curing accelerator (average particle size  
4.5  $\mu\text{m}$ ) including DBU.

Epikote 828 (100 parts by weight, epoxy equivalent  
35 amount 190 g/eq.) and Rikacid MT-500 (91 parts by weight,

methylnetetrahydrophthalic anhydride, manufactured by New Japan Chemical Co. Ltd.) were mixed, 1 part by weight of the obtained microencapsulated curing accelerator was added and the mixture was uniformly mixed to give a curable composition.

5 The composition was evaluated in the same manner.

(Comparative Example 7): crystalline epoxy resin

In the same manner as in Example 5 and using 4,4'-glycidyl-3,5,3',5'-tetramethyl-biphenyl (165 parts by weight, epoxy equivalent amount 165 g/eq.), ethylenediamine (30 parts  
10 by weight, amine equivalent amount 30 g/eq) and methylethylketone (100 parts by weight), a curable composition was obtained and subjected to the evaluation in the same manner.

The evaluation results of Examples are shown in Table 1  
15 and the evaluation results of Comparative Examples are shown in Table 2.

In Examples 1-6, good preservation stability was shown with almost no progress of the reaction even after the lapse of 1000 hr at 50°C. At 150°C, rapid hardening occurred and the  
20 properties of the cured product were shown to be sufficiently suitable for use for formation of coated films and adhesion.

In contrast, in Comparative Examples 1-5, a curing reaction proceeded from immediately after mixing, and preservation at room temperature, not to mention at 50°C, was  
25 shown to be difficult.

The microencapsulated curing accelerator of Comparative Example 6 was clarified to show poor adaptability to a dispersing machine having high dispersion force such as a three roll mill. The crystalline epoxy of Comparative Example  
30 7 showed insufficient preservation stability.

Table 1

composition		Example 1	Example 2	Example 3	Example 4	Example 5	Example 6
epoxy compound		Epikote 828	Epikote 828	Epikote 828	EOCN-102S	Epikote 1001	Epikote 1004
multifunctional amine compound		(I)	(V)	(IX)	(I)	(I)	(I)
solvent		none	none	none	none	MEK	MEK
volume							
average particle size	$\mu\text{m}$	0.8	0.9	0.8	0.7	1.8	1.2
preservation stability		O	O	O	O	O	O
curability		O	O	O	O	O	O
pencil hardness		4H	4H	4H	5H	3H	3H
adhesive property	N/cm	7.1	7.4	8.2	7	6.5	6.8
water resistance		O	O	O	O	O	O



Table 2

composition	Comparative Example 1	Comparative Example 2	Comparative Example 3	Comparative Example 4	Comparative Example 5	Comparative Example 6	Comparative Example 7
epoxy compound	Epikote 828	Epikote 828	Epikote 828	Epikote 828	Epikote 828	Epikote 828	crystalline epoxy
multifunctional amine compound	o-phenylene diamine	m-phenylene diamine	p-phenylene diamine	ethylene diamine	diamino-diphenyl ether	acid anhydride, micro-encapsulated curing accelerator	ethylene diamine
	solvent	none	none	none	none	none	MEK
	volume average particle size $\mu\text{m}$	(dissolved)	(dissolved)	(dissolved)	(dissolved)	10	8
	preservation stability	X	X	X	X	$\Delta$	$\Delta$
curability	O	O	O	O	O	O	O
pencil hardness	4H	4H	4H	2H	3H	3H	3H
adhesive property	5.2	6.1	5.5	7.1	6.7	4.6	4.8
water resistance	X	X	X	X	X	O	O

### **Industrial Applicability**

As described above, the curable composition of the  
5 present invention is one-component, shows high preservation  
stability at normal temperatures, rapid curing reaction and  
superior properties of the cured product. It basically affords  
a simple form consisting of a base resin (epoxy compound) and  
a curing agent compound that reacts with the base resin at an  
10 equimolar ratio. The curable composition of the present  
invention is useful for adhesives, daubs, coating materials,  
ink, paints, resins for molding and the like.

This application is based on a patent application No.  
15 2003-418789 filed in Japan, the contents of which are  
incorporated in full herein by this reference.